

*Appl. No. 09/909,001**Rule 1.312 Amendment dated May 12, 2004**Case No. 4543 US***Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

We claim:

1. (currently amended) A method of isolating one strand of a double-stranded target nucleic acid, comprising: (i) contacting a double-stranded target nucleic acid comprising a first strand and a second strand with a competitor oligo capable of hybridizing to the first strand under conditions in which the first strand dissociates from the second strand and hybridizes with the competitor oligo to form a first-strand:competitor oligo ~~heterduplex~~ heteroduplex; and (ii) isolating the dissociated second strand.

2. (currently amended) A method of isolating one strand of a double-stranded target nucleic acid, comprising: (i) contacting a double-stranded target nucleic acid comprising a first strand and a second strand with a competitor oligo capable of hybridizing to the first strand under conditions in which the first strand dissociates from the second strand and hybridizes with the competitor oligo to form a first-strand:competitor oligo ~~heterduplex~~ heteroduplex; (ii) isolating the heteroduplex, and (iii) dissociating the ~~heteroduplex~~ heteroduplex and isolating the first strand.

3. (cancelled)

4. (cancelled)

5. (cancelled)

6. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the double-

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stranded target nucleic acid is a double-stranded DNA.

7. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the double-stranded target nucleic acid is a double-stranded DNA/RNA hybrid duplex.

8. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the competitor oligo is composed of between 7 and 40 nucleobases.

9. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the double-stranded target nucleic acid has the formula:

TAIL 1-SEQUENCE-TAIL 2 TAIL 1'-SEQUENCE'-TAIL 2' wherein: TAIL 1 represents a first tail nucleobase sequence; SEQUENCE represents a target nucleobase sequence; TAIL 2 represents a second tail nucleobase sequence; TAIL 1' represents a nucleobase sequence that is complementary to TAIL 1; SEQUENCE' represents a nucleobase sequence that is complementary to SEQUENCE; and TAIL 2' represents a nucleobase sequence that is complementary to TAIL 2.

10. (original) The method of claim 9 in which a portion of the competitor oligo is capable of hybridizing to TAIL 1 and another portion of the competitor oligo is capable of hybridizing to TAIL 2.

11. (original) The method of claim 9 in which a portion of the competitor oligo is capable of hybridizing to TAIL 1' and another portion of the competitor oligo is capable of hybridizing to TAIL 2'.

12. (original) The method of claim 9 in which TAIL 1 and TAIL 2 comprise non-standard synthetic nucleobases.

13. (original) The method of claim 9 in which TAIL 1 and TAIL 2 are not complementary to one another.

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14. (previously presented) The method of claim 1 or 2 or 27 or 28, in which the competitor oligo includes a capture moiety.

15. (original) The method of claim 14, in which the capture moiety is one member of a pair of molecules that specifically bind to each other.

16. (original) The method of claim 14, in which the capture moiety is biotin.

17. (previously presented) The method of 14, in which the capture moiety is a solid support.

18. (original) The method of claim 17, in which the solid support is magnetic.

19. (original) The method of claim 14 in which the capture moiety is a capture sequence.

20. (previously presented) The method of claim 14 in which the capture moiety is a charged group.

21. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the competitor oligo is capable of hybridizing to only the first or the second strand of the double-stranded target nucleic acid.

22. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the contacting step is carried out at a cationic strength in the range of 0 to 10 mM, a pH in the range of 6 to 8, and a temperature in the range of 20 to 40° C.

23. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the competitor oligo is a PNA and optionally includes from 1 to 4 positively charged nucleobase interlinkages.

24. (previously presented) The method of claim 1 or 2 or 27 or 28 in which the

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competitor oligo comprises nonstandard synthetic nucleobases.

25. (previously presented) A method of isolating one strand of a double-stranded target nucleic acid, comprising the steps of: (i) dissociating the double-stranded target nucleic acid into a first strand and a second strand; (ii) contacting the dissociated target nucleic acid with a competitor oligo capable of hybridizing to only the first strand under conditions which kinetically favor competitor oligo first-strand hybrid formation and kinetically disfavor reannealing of the first and second strands, said competitor oligo being conjugated with a moiety that facilitates capture of competitor oligo: first-strand hybrids; (iii) capturing the competitor oligo: first strand hybrid, and (iv) dissociating the heteroduplex and isolating the first strand.

26. (original) The method of claim 25 wherein the competitor oligo is a PNA.

27. (previously presented) A method of isolating one strand of a double-stranded target nucleic acid, comprising: (i) dissociating the double-stranded target nucleic acid into a first strand and a second strand; (ii) contacting the dissociated target nucleic acid with a competitor oligo capable of hybridizing to the first strand under conditions which favor first-strand: competitor oligo heteroduplex formation and disfavor reannealing of the first and second strands; and (iii) isolating the dissociated second strand.

28. (previously presented) A method of isolating one strand of a double-stranded target nucleic acid, comprising: (i) dissociating the double-stranded target nucleic acid into a first strand and a second strand; (ii) contacting the dissociated target nucleic acid with a competitor oligo capable of hybridizing to the first strand under conditions which favor first-strand: competitor oligo heteroduplex formation and disfavor reannealing of the first and second strands; (iii) isolating the heteroduplex, and (iv) dissociating the heteroduplex and isolating the first strand.